



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C08B 31/00, 15/00, 37/14, A61L 15/00	A1	(11) International Publication Number: WO 98/27117 (43) International Publication Date: 25 June 1998 (25.06.98)
(21) International Application Number: PCT/NL97/00708 (22) International Filing Date: 18 December 1997 (18.12.97) (30) Priority Data: 96203594.5 18 December 1996 (18.12.96) EP <i>(34) Countries for which the regional or international application was filed:</i> NL et al. 96203595.2 18 December 1996 (18.12.96) EP <i>(34) Countries for which the regional or international application was filed:</i> NL et al. (71) Applicant (for all designated States except US): NEDERLANDSE ORGANISATIE VOOR TOEGEPAST-NATUURWETENSCHAPPELIJK ON- DERZOEK (TNO) [NL/NL]; Schoemakerstgraat 97, NL-2628 VK Delft (NL). (72) Inventors; and (75) Inventors/Applicants (for US only): BESEMER, Arie, Cornelis [NL/NL]; H. v.d. Boschstraat 111, NL-3958 CC Ameron- gen (NL). THORNTON, Jeffrey, W. [US/NL]; Singel 18, NL-1402 NT Bussum (NL).		(74) Agent: DE BRUIJN, Leendert, C.; Nederlandsch Octrooi- bureau, Scheveningseweg 82, P.O. Box 29720, NL-2502 LS The Hague (NL). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the</i> <i>claims and to be republished in the event of the receipt of</i> <i>amendments.</i>
(54) Title: SUPERABSORBENT MATERIAL AND METHOD FOR PRODUCING SAID MATERIAL (57) Abstract A superabsorbent polysaccharide derivative is obtained by oxidation of the polysaccharide to the extent that the derivative contains 0.2–3.0 carboxyl group per monosaccharide unit and cross-linking the oxidised polysaccharide with 0.001–0.2 equivalent of cross-linking agent per monosaccharide unit. The polysaccharide may also be cross-linked prior to oxidation.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

Superabsorbent material and method for producing said material

The present invention relates to a superabsorbent material based on polysaccharides and to a method of producing such material. A superabsorbent material is defined as a material capable of absorbing at least 20 times its own weight of water.

5 Superabsorbent materials of various types are known in the art. Examples are crosslinked polyacrylates and polysaccharides grafted with polyacrylates. A problem related to the use of such superabsorbent materials is that such materials are normally based on non-renewable and/or non-biodegradable raw materials. Consequently, there is a need for superabsorbent materials which are wholly based on renewable raw materials,
10 such as polysaccharides, and are degraded by microorganisms or other natural agents after being disposed of.

EP-A-489424 discloses modified starch compositions useful as absorbable dusting powders, obtained by slight hypochlorite oxidation of starch resulting in the presence of 0.05 - 0.5 wt.% of carboxyl groups (= 0.002 - 0.018 carboxyl groups per
15 monosaccharide unit), followed by partial crosslinking with 0.05 - 1.1 wt.% of phosphorus oxychloride. The powders obtained according to this prior art are not suitable for use in absorbent articles such as diapers, sanitary napkins and the like due to the fact that they do not swell to a significant extent. Kulicke et al (*Starch/Stärke*, **41** (1989) 140-146; *ibid.* **42** (1990) 134-141) have studied the effect of crosslinking with epichlorohydrin
20 and sodium trimetaphosphate, respectively, on the absorption capacity of starch, amylopectin and commercial oxidised starch, which contains carbonyl (aldehyde or ketone) groups and carboxylic acid groups. The degree of oxidation of such oxidised starch is presumably lower than 10%. The purpose of using oxidised starch was to obtain thin boiling starch (lower molecular weight). According to these authors, the absorption
25 capacity decreases with increased crosslinker concentration and the absorption capacity of crosslinked oxidised starch is higher than that of crosslinked native starch. Although a considerable swelling capacity in pure water was reported, the swelling capacity of the crosslinked polymers in water containing sodium chloride or other salts drops to about 15 g per g polymer. US-2,929,811 discloses crosslinking (using epichlorohydrin) and
30 subsequent slight oxidation (< 5 mol%), using periodate and chlorite, of starch for improving the viscosity characteristics of the starch. No water-absorbing properties are reported.

EP-A-23561 discloses a water-absorbing crosslinked polycarboxycellulose which is produced by carboxymethylation of cellulose, crosslinking by acid-catalysed internal esterification, and slight oxidation (< 5 mol%) using periodate/bromine, lead tetraacetate or hydrogen peroxide. A disadvantage of carboxymethyl celluloses is that they are relatively expensive.

It has now been found that a superabsorbent material having an absorption capacity of at least 20 times its own weight, and based on natural, biodegradable polysaccharides, can be obtained by introducing at least 0.2 carboxyl groups per monosaccharide unit, at least 0.1 of which is formed by oxidising a carbinol group of the polysaccharide, and subsequently slightly crosslinking the oxidised polysaccharide.

The polysaccharides to be used according to the present invention are in particular α -glucans like starch, amylose and amylopectin, β -glucans like cellulose, galactomannans like guar gum (guaran) and locust bean gum, glucomannans including e.g. xanthan gum, fructans, (arabino)xylans, galactans including alginates and pectins, as well as non-ionic derivatives such as hydroxyethyl and hydroxypropyl derivatives of such polysaccharides. Starch and guar, and to a somewhat lesser extent, cellulose, are preferred for economic reasons. The chain length of the polysaccharides is important although there is no critical minimum for the molecular weight. In general, polysaccharides having a molecular weight of more than 1,000 are preferred. A molecular weight above about 25,000 may have a positive effect on the properties of the oxidised product.

The polysaccharides to be used according to the present invention may also be carboxymethylated or carboxyethylated, especially in the case of α -glucans like starch, galactomannans and glucomannans. Other carboxyalkylated polysaccharides include the half esters obtained from cyclic anhydrides such as succinic and maleic anhydride (groups having the formula $-O-CO-CH_2-CH_2-COOH$ or $-O-CO-CH=CH-COOH$), and addition products of maleic half esters to which sulphite has been added (groups having the formula $-O-CO-CH_2-CH(SO_3H)-COOH$). The degree of carboxyalkylation is preferably between 0 and 1.5, in particular between 0.1 and 1.0 carboxyalkyl groups per monosaccharide unit. Carboxymethylation and carboxyethylation can be performed in a conventional manner, i.e. by reaction of the (oxidised or non-oxidised) polysaccharide with monochloroacetate, or acrylonitrile followed by hydrolysis, respectively, or by hydroxyethylation (or hydroxypropylation) followed by oxidation of the primary hydroxy groups, e.g. using a nitroxyl catalyst as described below. Thus, a useful type of products according to the

invention containing both 6-carboxyl groups and carboxymethyl groups can be obtained in several ways: oxidation of carboxymethylated polysaccharide, carboxymethylation of 6-oxidised polysaccharide, (primary hydroxyl) oxidation of hydroxyethylated polysaccharide, and hydroxyethylation and oxidation of 6-oxidised polysaccharide.

5 The oxidation of the polysaccharide is an essential step of the present invention. The oxidation should be performed to a substantial degree, i.e. until at least 0.1 of the carbinol group per monosaccharide unit has been oxidised to a carboxyl group. The term carbinol group covers both primary, exocyclic hydroxymethyl groups ($-\text{CH}_2\text{OH}$) and secondary, usually endocyclic hydroxymethylene groups ($-\text{CHOH}-$) of the polysaccharide.
10 Preferably at least 0.15 or even at least 0.2 carbinol group have been oxidised to a carboxyl group per monosaccharide unit. Additional carboxyl groups may be present in the form of carboxymethyl groups introduced by substitution on a hydroxyl group .

 Oxidation of polysaccharides is well-documented. Oxidation can be performed using various oxidising agents, which results in various degrees of oxidation, various
15 degrees of polymerisation and different sites of oxidation. Oxidation of polysaccharides can be focused at primary hydroxyl groups, like the 6-hydroxyl group in the anhydro-glucose units of glucans, which results in carboxyl-polysaccharides with preserved ring structures. On the other hand, the oxidation can be mainly directed at a vicinal diol function present in the monosaccharide rings, such as the C2-C3 site in anhydroglucose
20 units. This results in cleavage of the monosaccharide units with the production of dialdehyde and/or dicarboxyl functions.

 As an example, the oxidation of starch with nitrite and nitrate in phosphoric acid, mainly resulting in 6-carboxy starch, is described in NL patent application 9301172. An improved oxidation of the 6-hydroxyl groups in starch, using a hypohalite in the presence
25 of a di-tert-alkylnitroxyl catalyst, is disclosed in WO 95/07303. Examples of oxidation of glucans at the C2-C3 function include the process according to EP-A-427349, using low levels of hypobromite, and the process according to WO 94/21690, which uses hydrogen peroxide in the presence of alkali metals or transition metals. WO 95/12619 describes an improved oxidation of starch with periodic acid, resulting in dialdehyde starch
30 with extensive regeneration of a periodic acid. The dialdehyde starch can be further oxidised to dicarboxyl starch using e.g. sodium chlorite and/or hydrogen peroxide. Also, dialdehyde starch can be further oxidised with iodine or bromine or with nitrogen dioxide producing dicarboxy or up to tricarboxy starch. Other known oxidation methods include

metal-catalysed oxidation, e.g. using ruthenium, anhydrous oxidation using nitrogen dioxide in e.g. halocarbons and enzymatic and chemo-enzymatic oxidation of starch, guar and other polysaccharides, and these can also be used in the present invention. In case of galactomannans, which have terminal galactose units, enzymatic oxidation using galactose oxidase (EC 1.1.3.9) can also be used to introduce aldehyde groups, which can easily be converted to carboxyl groups, e.g. using hypiodite.

Any of the known oxidation processes can be used according to the present invention. Also, a combination of oxidation processes and/or oxidation agents can be used. Oxidation of primary hydroxyl functions in the polysaccharides has a slight advantage, in that the resulting carboxyl functions are on average at a wider distance than the carboxyl groups introduced by C2-C3 oxidation and therefore allow a somewhat lower average degree of oxidation. The oxidation of starch and other polysaccharides using hypochlorite in the presence of di-tert-alkylnitroxyl catalysts, in particular tetramethylpiperidine-N-oxyl (TEMPO) is particularly efficient.

The degree of oxidation is such that more than 1.0 out of each 10 monosaccharide units is oxidised to produce a carboxy- or dicarboxy-monosaccharide unit. The number of dicarboxy-monosaccharide units may be up to 10 out of 10, but is preferably between 2 and 8 out of 10, and most preferably between 2 and 6. Expressing the degree of oxidation in a carboxyl content, the polysaccharide contains on average 0.2-3.0 carboxyl group per monosaccharide unit, preferably 0.4-1.6 and most preferably 0.4-1.2 carboxyl group per monosaccharide unit. The total carboxyl content (both resulting from carboxy-alkylation and from oxidation) in case of carboxyalkylated polysaccharides is preferably between 0.2 and 2 carboxyl groups per C-6/7 unit.

Dicarboxy-monosaccharide units especially result from C2-C3 oxidation such as hypohalite oxidation. The dicarboxy-monosaccharide units are usually ring-opened units having a higher charge distribution and more water-binding capacity. On the other hand C2-C3 oxidation leads to more extensive degradation (depolymerisation) of the polysaccharide unless more expensive oxidising agents such as periodate are used. Mono-carboxy-monosaccharide units especially result from C6 oxidation such as TEMPO oxidation. Such oxidations are usually less accompanied by degradation.

The oxidised polysaccharide is subsequently reacted with a crosslinking agent. Crosslinking agents are reagents containing two or more functions capable of reacting with a hydroxyl group, resulting in intra- and inter-molecular bonds between different mono-

saccharide units. Suitable crosslinking agents may act on the hydroxyl groups of different polysaccharide chains and include divinyl sulphone, epichlorohydrin, diepoxybutane, diglycidyl ethers, diisocyanates, cyanuric chloride, trimetaphosphates, phosphoryl chloride, and mixed anhydrides, and also inorganic crosslinkers such as aluminium and zirconium ions, but are not restricted to these examples. Crosslinking can also be performed using carboxyl or aldehyde groups formed by oxidation or carboxyl groups introduced by carboxyalkylation, e.g. using polyols, polyamines or other polyfunctional reagents. Esterification and other crosslinking methods described herein can also be effected intramolecularly at the surface between the carboxyl group of one polysaccharide chain and a hydroxyl group of another chain as known in the art. This inter-chain crosslinking can be catalysed by an acid or a multivalent ion such as magnesium or calcium, or by heating. Crosslinking of starch and other polysaccharides is well-known in the art. A description of crosslinking agents and reaction conditions can be found e.g. in *"Starch Derivatives: Production and Uses"* by M.W. Rutenberg and D. Solarek, Acad. Press Inc., 1984, pages 324-332.

The crosslinking after oxidation to the desired level of crosslinking can be achieved by reacting the oxidised polysaccharide with 0.1-20 mol.% of crosslinking agent, preferably 0.2-10 mol.% and most preferably 0.3-2 mol.%. Optimum crosslinking rates and conditions depend on the particular polysaccharide and on the particular crosslinking agent, as well as on the type of oxidation: for C6 oxidation, the optimum rate may be e.g. 0.3-0.5 mol%, while for C2-C3 oxidation the optimum may be somewhat higher, e.g. 0.5-2 mol%. Reagents such as diglycidyl ether are usually reacted under acidic conditions, whereas most other reagents are reacted under neutral or alkaline conditions. Reaction conditions and working-up procedures are described e.g. in US 4,582,865. Work-up may be done by drawing off the aqueous solution of crosslinking agent from the crosslinked or gelatinised polysaccharide, drying the latter under air, under reduced pressure and/or in a drying oven, optionally using water-miscible and/or volatile solvents such as methanol, ethanol, isopropanol, acetone, dioxane or tetrahydrofuran, followed e.g. by freeze-drying, spray-drying, vacuum-drying or other appropriate technique. Crosslinking preferably results in a gel, that may precipitate from the reaction medium.

It is sometimes advantageous to perform a crosslinking before the oxidation reaction. This crosslinking can be performed using the same agents and the same conditions as the post-crosslinking described above. In general, the level of prior

crosslinking is lower than that of the post-crosslinking. For example, the prior crosslinking can be performed using 0.05–1.0, preferably 0.1–0.8, most preferably 0.1–0.5 mol.% of crosslinking agent, such as epichlorohydrin. It is preferred that the salt concentration of the polysaccharide solution during crosslinking is low; this results in a better accessibility of the polysaccharide for the crosslinking agents. Some pre-crosslinked starches are commercially available and can be used in the present invention. In some instances, prior crosslinking, without post-crosslinking, may be sufficient, e.g. when oxidation of the polysaccharide is performed at the 6-position, such as with TEMPO, or when oxidation is combined with carboxyalkylation.

The superabsorbent polysaccharide according to the invention has improved absorption characteristics, especially of body fluids which contain various salts and non-ionic substances. The product is particularly suitable for the production of absorbent hygiene articles, such as diapers, sanitary napkins and the like. Such articles can be produced entirely on the basis of the polysaccharides according to the invention, but they can also contain conventional absorbent materials, such as cellulose pulp in addition to the absorbents according to the invention.

Example 1: Preparation of 2,3-dicarboxy starch

In a cooled dispersion of non-crosslinked potato starch (16.2 grams of dry substance) in 300 ml water sodium periodate was added at 5°C (7.5 g, 9.6 g and 12.8 g to obtain a degree of oxidation of 35, 45 and 60%, respectively). The pH was adjusted to 5 and the mixture was stirred in the dark for 20 hours at 5°C. Unreacted sodium iodate was then removed by filtration and repeated washings with water.

The dialdehyde starch thus obtained (15.6 g) was suspended in chilled water (100 ml) and the pH was adjusted to 5. Stoichiometric amounts of sodium chlorite and hydrogen peroxide were added, i.e. 2 moles of NaClO₂ (8.0, 10.2 and 13.6 g, respectively) and 2 moles of H₂O₂ (7.3, 9.3 and 12.4 g, respectively) were used per mole of sodium periodate. The mixture was stirred at 5°C, keeping the pH at 5 by addition of NaOH. Addition of water was required to keep the mixture stirred. After three hours the pH remained more or less constant without the need of adding alkali. After 20 hours, the reaction mixture was poured into three volumes of ethanol. A gelly precipitate was formed, from which the aqueous solution was decanted. The precipitate was dissolved in water and the procedure was repeated. The second precipitate was a solid and was collected by filtration and dried

in vacuo at 50°C. The overall yield was 90%, calculated on the starting starch, and the carboxyl content was 32, 42 and 55 %, respectively.

Example 2: Preparation of pre-crosslinked 2,3-dicarboxy starch

The procedure of example 1 was repeated, starting with epichlorohydrin-crosslinked starch
5 instead of native starch. The overall yield and the carboxyl content were the same.

Example 3: Crosslinking of 2,3-dicarboxy starch

An amount of 15, 20 or 30 wt.% of 45% oxidised starch (dicarboxy starch, example 1) was dissolved in 100 ml of 0.1 M aqueous NaOH for 4 hours. At room temperature an amount of 2, 5 or 8 mole % of divinyl sulphone (0.5 mole % being 1.8/200 g DVS per
10 g of dicarboxy starch) was added to the solution and mixed. The reaction mixture was left overnight at room temperature. The resulting gel was then washed with distilled water and subsequently with acetone and then dried at room temperature. The absorption capacity of the resulting crosslinked oxidised starch was measured for pure water and for synthetic urine. The composition of the synthetic urine (SU) is as follows:

15 300 mM of urea

60 mM of KCl

130 mM of NaCl

3.5 mM of MgSO_4

2.0 mM of $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$

20 1 g/l of a 0.1% solution of Triton X-100 (Riedel de Haen); in deionised water.

The results are given in table 1.

Example 4: Crosslinking of pre-crosslinked 2,3 dicarboxy starch

The procedure of example 3 was repeated, using pre-crosslinked, oxidised starch according to example 2. The results are given in table 1.

25 **Example 5: Crosslinked dicarboxy starch**

Following the procedure of examples 1 and 3, starch was oxidised with periodate and chlorite to an oxidation degree of 50% and subsequently crosslinked as a 30 wt.% solution with 2 mol% divinyl sulphone. The product was isolated by precipitation from methanol and had a free swelling capacity (FSC) in synthetic urine (see example 3) of
30 42 g/g.

Table 1
Water absorption capacities of crosslinked 45% oxidised dicarboxy-starch

	% pre-XL / wt.% sol / % post-XL	abs. capac. 60 min.	abs. capac. 120 min.	retention capac.
5	0 / 20 / 2	25.9	26.2	19.7
	0 / 30 / 2	22.9	20.8	15.7
	0 / 20 / 5	25.0	24.8	18.7
	0 / 15 / 8	24.2	22.9	17.4
	0 / 20 / 8	21.5	22.1	14.5
10	0.5 / 30 / 2	29.1	28.9	16.7
	0.5 / 20 / 5	24.4	25.6	16.2
	0.5 / 30 / 5	21.6	20.4	14.4
	0.5 / 15 / 8	26.4	25.9	15.3
	0.5 / 20 / 8	24.4	24.2	17.4

- 15 % pre-XL mole percentage of epichlorohydrin used for pre-crosslinking with
 respect to anhydroglucose units
 wt.% sol. weight percentage of oxidised starch before post-crosslinking
 % post-XL mole percentage of divinyl sulphone used for post-crosslinking with
 respect to anhydroglucose units

20 **Example 6: Preparation of 6-carboxy starch**

A dispersion of 16.2 g (dry matter) of non-crosslinked potato starch (100 mmol of monosaccharide units) in 300 ml of water was prepared. 45 mg of TEMPO and 450 mg of sodium bromide were dissolved in the gelatinised mixture. A solution of sodium hypochlorite, 3.5 meq of hypochlorite per ml, was added (90, 146 and 192 ml in three experiments to obtain a degree of oxidation of 40, 60 and 80%, respectively). The pH of the mixture was kept at 10 throughout the process by adding sodium hydroxide solution using a pH stat. After a fast reaction, the reaction ceased within 30 minutes, indicated by the fact that no further sodium hydroxide addition occurred. The degree of oxidation was determined from the consumption of hydroxide. After 1 hour, 100 mg of sodium borohydride was added to remove free aldehyde groups. The product was isolated by pouring the reaction mixture into ethanol as described in example 1. The yield was 90%, and the carboxyl content corresponded to the degree of oxidation.

Example 7: Crosslinking of 6-carboxy starch

6-Carboxy starch (degree of oxidation 40% or 60%, example 6) was dissolved in water (concentration 20%) at pH 11-12 and cooled to 0-2°C. A 10% solution of divinyl sulphone was added (2 or 3 mole % = 8 or 12 mg, respectively) and after homogenisation the mixture was allowed to stand for 4 hours. The temperature was allowed to rise to 20°C. The crosslinked polymer was isolated by pouring the mixture in ethanol (1 volume of reaction mixture to 3 volumes of ethanol). The precipitate was collected by filtration, washed twice with absolute ethanol and dried in vacuo. The water binding capacity after 15 minutes using synthetic urine was determined to be:

40% oxidation	2 mole% of water	23 g/g
40% oxidation	3 mole% of water	23 g/g
60% oxidation	2 mole% of water	21 g/g
60% oxidation	3 mole% of water	21 g/g

After 2 hours, the water binding capacity rises with another 10-15 g/g.

Example 8: Crosslinked 6-carboxy starch

Following the procedure of examples 6 and 7, starch was oxidised at the 6-position using an amount of oxidising agent to obtain a degree of oxidation of 60%, and was subsequently crosslinked with divinyl sulphone in an amount of 0.5 mol%. After work-up the product had an FSC (free swelling capacity) in synthetic urine (see example 3) after 2 hours of 53 g/g. The FSC in 0.9 % NaCl was 62 g/g. When the procedure was repeated started with starch which was pre-crosslinked with 0.05 wt.% epichlorohydrin, an FSC in synthetic urine of 49 g/g was obtained.

Example 9: Pre-crosslinking and 6-oxidation of starch

Potato starch (5.0 g) was dissolved in 0.5 mol/l sodium hydroxide. Then 2.5 g of sodium trimetaphosphate (26 mol% with respect to starch) was added. After mixing thoroughly, the samples were kept at 40°C for 2 hours and then at room temperature for 16 hours. The gels which were obtained were broken into small particles using a mechanical stirrer. The particles were washed with demineralised water until the pH was below 9. The crosslinked starch was then oxidised with TEMPO and hypochlorite/bromide (see example 6), using an amount of hypochlorite corresponding to a degree of oxidation of 60%. The product was then precipitated with ethanol, washed with acetone and dried in a vacuum oven. The

FSC value after 1 hour in synthetic urine was 26.7 g/g.

Table 2

FSC and CRC (centrifuge retention capacity) values of post-crosslinked oxidised carboxymethyl starch; values corrected for water still present in sample

5	% oxidation	mol% DVS	FSC (g/g) SU/2h	FSC (g/g) 0.9% NaCl/2h	CRC (g/g) SU
	17.5	0.1	22.4	31.5	1.6
	17.5	0.2	49.6	77.8	10.2
	17.5	0.3	76.6	87.9	23.5
	17.5	0.4	61.1	80.2	17.6
10	17.5	0.5	61.2	79.9	16.7
	35	0.2	64.9	—	21.2
	35	0.4	55.2	—	35.2
	35	0.6	44.9	—	30.8
	40	0.2	61.1	—	23.2
15	40	0.4	42.8	—	30.6
	40	0.6	30.8	—	20.4

Example 10: 6-Oxidation and crosslinking of carboxymethyl starch

Carboxymethyl starch, 10 g (degree of substitution 0.5) was dissolved in 800 ml of demineralised water. 40 mg of TEMPO and 1 g of NaBr were added. An amount of sodium hypochlorite was added corresponding to a degree of oxidation of 17.5 %. The hypochlorite was added in portions of 2.5 ml at a time. The pH was kept at 9.5 using 0.5 mol/l sodium hydroxide. After the oxidation reaction was complete, residual aldehyde groups were removed by adding 0.5 g of sodium borohydride and stirring for 1 hour. The product was then precipitated using 4 times the volume of ethanol, washed with acetone and dried in a vacuum oven.

After drying, the product (1 g) was crosslinked with divinyl sulphone (DVS). Different amounts of DVS were used: in mol% with respect to the starch, 0.05, 0.1, 0.2, 0.3, 0.4 or 0.5 mol% was used. The required amount of DVS was added as a 0.1 % solution to the samples which were dissolved in 4 ml 0.5 mol/l sodium hydroxide at 5°C. The

mixtures were vortexed strongly and placed overnight at 5°C. The next day the gels were broken with a mechanical stirrer. Then 3 ml of 0.2 mol/l acetic acid and 17 ml of demineralised water was added. The gels were left swelling for 1 hour and then precipitated with ethanol, washed with acetone and dried in a vacuum oven. After the samples were dried, the FSC values were determined in synthetic urine (see example 3) or in 0.9% NaCl after 2 hours. The results are given in table 2.

The procedure of this example was repeated using an amount of hypochlorite equivalent to 35% and 40% of oxidation instead of 17.5%. The results are also given in table 2.

Example 11: Oxidation and crosslinking of guar

Purified guar gum, (1 g, 6.2 mmol monosaccharide units) was dissolved in 400 ml water and the solution was heated and stirred overnight. After cooling to 5°C 10 mg TEMPO and 100 mg NaBr were added. Oxidation was started by adding 3.0 ml of NaOCl solution in small steps. The pH was kept between 9.0 and 9.5 with 0.5 NaOH. After the reaction was completed, NaBH₄ was added and after 15 min. the pH of the solution was adjusted to 6.5 with 4 M HCl. The oxidised guar product was isolated by precipitating in 3 l of a 80/20 (v/v) mixture of ethanol/water and washing the precipitate with 1 l of ethanol (96%). The carboxyl content was determined using the method of Blumenkrantz (N. Blumenkrantz and G. Asboe-Hansen, *Anal. Biochem.* **54** (1973) 484-489): the oxidation degree was 35%.

The oxidised material was crosslinked with 3 mol of divinyl sulphone and a weak gel was obtained. The crosslinked product was isolated by precipitating the weak gel in ethanol. The precipitate was dried in a vacuum oven. The free swelling capacity (FSC) of the isolated product in 2 hours and the centrifugal retention capacity (CRC) were determined in synthetic urine (see example 3). The FSC was 40 g/g and the CRC was 25 g/g.

Claims

1. Superabsorbent polysaccharide derivative obtained by oxidation and crosslinking of a polysaccharide, characterised in that at least 0.1 carbinol group per monosaccharide unit of the polysaccharide derivative has been oxidised to a carboxyl group, the total number of carboxyl groups per monosaccharide unit being 0.2–3.0, and the derivative results from reaction with at least 0.001 equivalent of crosslinking agent per monosaccharide unit.
2. Superabsorbent polysaccharide derivative according to claim 1, containing 0.4–1.6 carboxyl group per monosaccharide unit, at least 0.2 of which is formed by oxidation of a carbinol group.
3. Superabsorbent polysaccharide derivative according to claim 1 or 2, wherein said polysaccharide derivative is a glucan, galactomannan or glucomannan, in particular starch or guar or a starch or guar derivative.
4. Superabsorbent polysaccharide derivative according to any one of claims 1–3, wherein 0.1–1 carboxyl groups per monosaccharide unit are present as carboxyalkyl groups.
5. Process of producing a superabsorbent polysaccharide derivative by oxidation and crosslinking a polysaccharide, characterised in that the polysaccharide is oxidised to obtain a polysaccharide containing an average of at least 0.2 carboxyl groups per monosaccharide unit and is subsequently reacted with 0.001–0.2 equivalent of a crosslinking agent.
6. Process according to claim 5, wherein the polysaccharide is oxidised using hypochlorite, optionally in the presence of a di-tert-alkylnitroxide.
7. Process according to claim 5, wherein the polysaccharide is oxidised using periodate, optionally followed by chlorite oxidation.
8. Process according to any one of claims 5–7, wherein the oxidised polysaccharide is reacted with 0.002–0.1, preferably 0.005 – 0.05 equivalents of crosslinking agent per monosaccharide unit.

9. Process according to any one of claims 5-8, wherein, before the oxidation, the polysaccharide is reacted with a crosslinking agent.
10. Process according to any one of claims 5-9, wherein, before or after the oxidation, the polysaccharide is carboxyalkylated to a degree of substitution of 0.1-1 carboxyalkyl groups per monosaccharide unit.
11. Process of producing a superabsorbent polysaccharide derivative by oxidation and crosslinking a polysaccharide, characterised in that the polysaccharide is crosslinked by reaction with a crosslinking agent and subsequently oxidised using hypochlorite in the presence of a di-tert-alkylnitroxide to a degree corresponding to an average of at least 0.2 carboxyl groups per monosaccharide unit.
12. Process of producing a superabsorbent polysaccharide derivative by oxidation and crosslinking a polysaccharide, characterised in that the polysaccharide is an α -glucan, galactomannan or glucomannan which is carboxyalkylated to a degree of 0.1-1 carboxyalkyl groups per monosaccharide unit and oxidised to a degree corresponding to at least 0.1 carboxyl groups per monosaccharide unit, the total average number of carboxyl group per monosaccharide unit being at least 0.2, and which is pre- and/or post-crosslinked by reaction with a crosslinking agent or by interchain esterification.
13. Process according to any one of claims 5-12, wherein the crosslinking agent is selected from epichlorohydrin, a trimetaphosphate, phosphoryl chloride, divinyl sulphone, diglycidyl ether, a diisocyanate, or a mixed anhydride.
14. Process according to any one of claims 5-13, wherein the polysaccharide is an α -glucan, galactomannan or glucomannan.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/NL 97/00708

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C08B31/00 C08B15/00 C08B37/14 A61L15/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C08B A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 023 561 A (MONTEDISON S.P.A.) 11 February 1981 cited in the application see page 4, line 6 - line 14	1-14
Y	CHEMICAL ABSTRACTS, vol. 97, no. 6, 9 August 1982 Columbus, Ohio, US; abstract no. 40606, "Paper strenghtening agent" XP002060693 see abstract & DATABASE WPI Week 8213 Derwent Publications Ltd., London, GB; AN 24766E see abstract	1-14

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

30 March 1998

Date of mailing of the international search report

16/04/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax (+31-70) 340-3016

Authorized officer

Lensen, H

INTERNATIONAL SEARCH REPORT

Int. J. Application No.

PCT/NL 97/00708

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5 346 892 A (FITT ET AL.) 13 September 1994 see column 1, line 18 - line 22 ---	1-14
A	US 2 989 521 A (F.R.SENTI ET AL.) 20 June 1961 cited in the application ---	
A	US 2 929 811 A (B.T.HOFREITER ET AL.) 22 March 1960 cited in the application -----	

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/NL 97/00708

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 23561 A	11-02-81	JP 1612608 C JP 2027361 B JP 56008402 A US 4340731 A	30-07-91 15-06-90 28-01-81 20-07-82
US 5346892 A	13-09-94	US 5126334 A BR 9301423 A CA 2093058 A JP 1993921 C JP 6239901 A JP 7023401 B US 5385608 A AT 137506 T AU 634947 B AU 8837191 A CA 2056939 A,C CN 1063115 A DE 69119205 D DE 69119205 T EP 0489424 A ES 2086463 T FI 915722 A HK 166196 A JP 2110722 C JP 4293902 A JP 8016122 B MX 9102354 A	30-06-92 05-10-93 04-10-93 22-11-95 30-08-94 15-03-95 31-01-95 15-05-96 04-03-93 11-06-92 06-06-92 29-07-92 05-06-96 14-08-96 10-06-92 01-07-96 06-06-92 13-09-96 21-11-96 19-10-92 21-02-96 01-06-92
US 2989521 A	20-06-61	NONE	
US 2929811 A	22-03-60	NONE	